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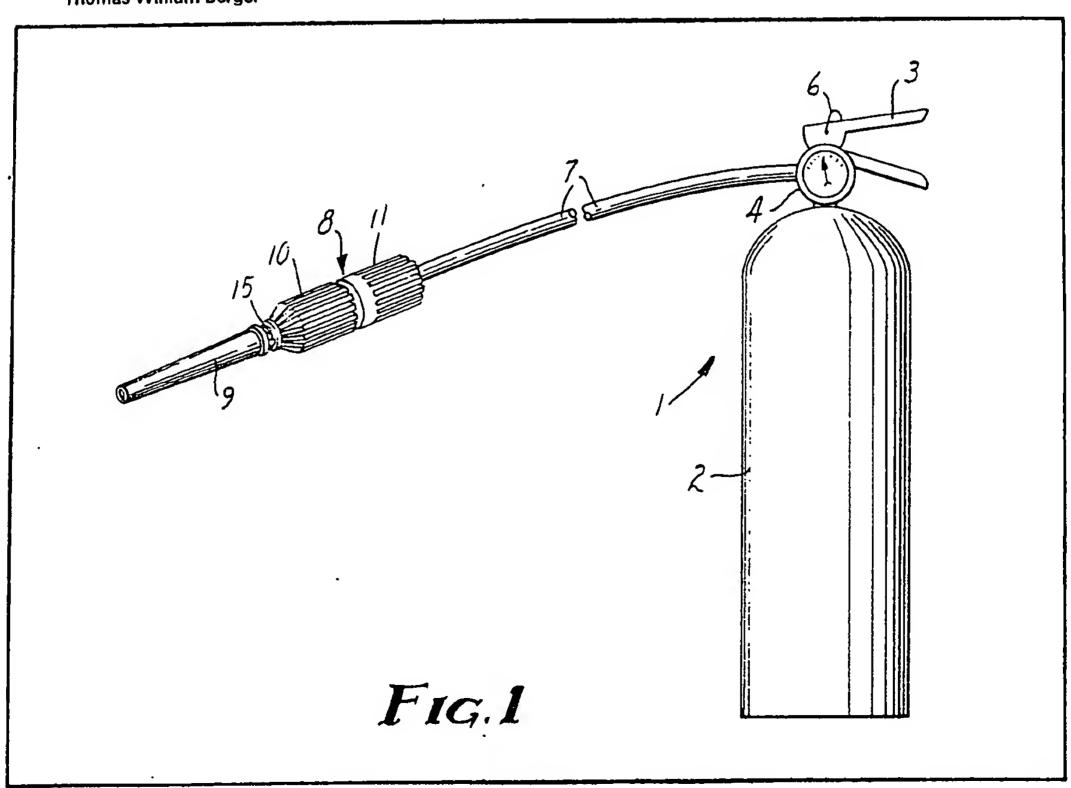
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(54) Aqueous Film-forming Foam Fire Extinguisher

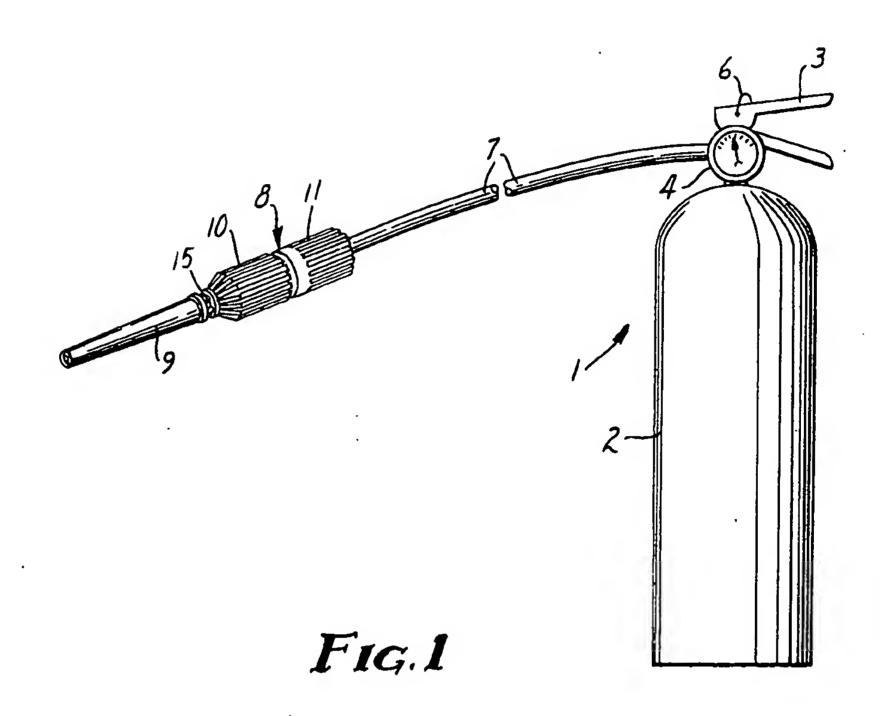
(57) A hand portable fire extinguisher comprising a tank adapted to contain pressurized water, a valve surmounting said tank, a hose connected to said valve, a cartridge holder connected to said hose, an air aspirating nozzle connected to said cartridge holder, and a cartridge loaded in said cartridge holder and

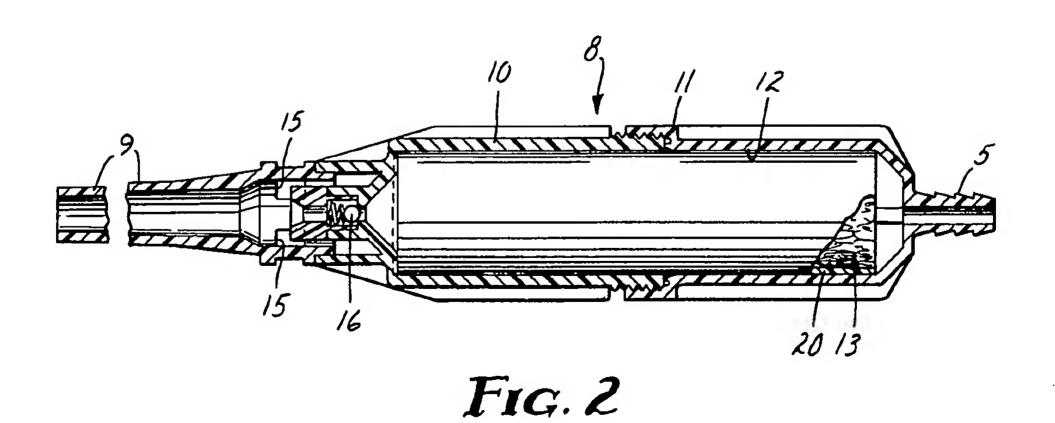
comprising a shaped body comprising a solid, coalesced mixture of water soluble fluoroaliphatic surfactant and water soluble fluorine-free surfactant, said shaped body having at least one exposed surface adapted to be contacted with water flowing through said cartridge holder to dissolve said shaped body and form an aqueous film-forming foam solution of relatively constant composition over the period of discharge of said water from said tank. Solid coalesced mixtures of water soluble fluorinated surfactant and water soluble fluorinefree surfactant, shaped bodies comprising such mixtures and cartridges containing a shaped body for use in the fire extinguisher.



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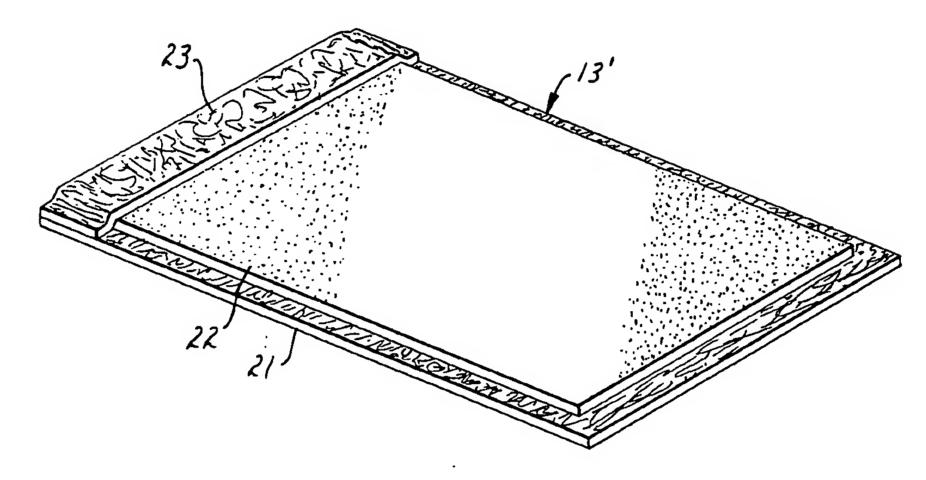


Fig.3

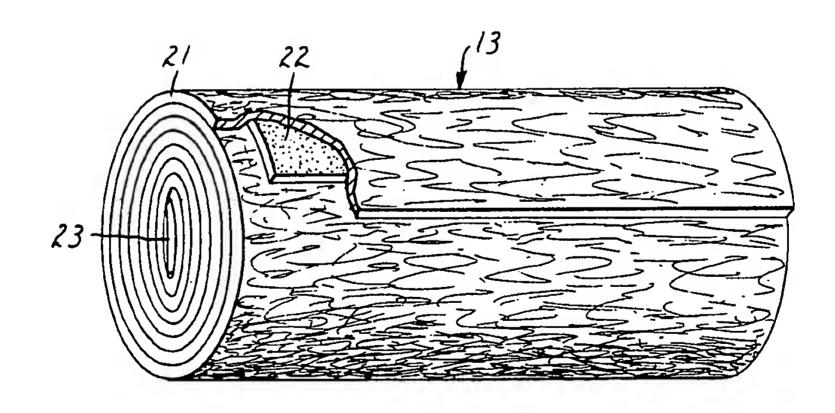


FIG. 4

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SPECIFICATION Agu us Film-forming Foam Fir Extinguisher

This invention relates to a hand portable fire extinguisher for discharging aqueous film-forming foam. In another aspect, it relates to a mixture of water soluble fluoroaliphatic surfactant and water soluble fluorine-free surfactant, and to shaped articles of said mixture. In a still further aspect, it relates to a method of extinguishing a fire of flammable liquid, or prevention of such fire, by applying to the surface of said liquid a foam of an aqueous film-forming solution from a hand portable fire extinguisher.

One of the most effective foam agents for extinguishing flammable liquid fires, such as fuel fires, is aqueous film-forming solution which is applied to fires as a foam, (such agents commonly 10 abbreviated as "AFFF"), a commercial fire extinguishing agent of this type being that sold as an aqueous liquid concentrate under the registered trademark "Light Water". This agent upon dilution with water has been used successfully to extinguish a host of test fires and such actual fires as a petrochemical storage facility fire, an oil tanker fire which had burned for three days, aircraft fires, and

numerous industrial fires of spilled fuel and solvent.

Surfactant compositions useful in or as fire fighting agents, including those of AFFF type, are 15 disclosed, for example, in U.S. Pat. Nos. 3,258,423 (Tuve et al), 3,562,156 (Francen), 3,661,776 (Fletcher et al), 3, 772,195 (Francen), 3,957,658 (Chiesa et al), 4,090,967 (Falk) and 4,149,599 (Chiesa). These compositions comprise solutions of water soluble, fluoroaliphatic surfactant, water soluble, fluorine-free surfactant, and water. Upon application of these compositions with, for example, 20 foam or water/fog equipment, a foam is generated which spreads over and floats on the surface of burning liquid, such as gasoline, forming a vapor-sealing film which extinguishes the fire; the film also secures non-ignited areas and prevents ignition. These compositions are usually provided and stored as aqueous concentrates which are diluted further with water upon use or they can be stored in the diluted form. Typical AFFF fire extinguishing systems are used for fire prevention and control of 25 relatively large hazardous or potentially hazardous flammable liquid bodies and require specially designed proportioning and delivery equipment which function by mixing the requisite amount of concentrate with water to produce the foam.

Recently, hand portable AFFF fire extinguishers have become commercially available, such as the 3M brand "Spoiler" fire extinguisher, described in 3M's bulletin Y-FEBIR(1062)MP, which can be 30 deployed and used to combat relatively small bodies of flammable liquids, such as might be common with automobile garages, paint shops, etc. These fire extinguishers are specially designed and contain about 2.5 gallons (9.5 liters) of premixed, ready-to-use, AFFF agent and are useful in combating small

"Class B" as well as "Class A" fires.

Briefly, according to one embodiment of this invention, there is provided a hand portable fire 35 extinguisher comprising a tank loaded with water under pressure, a squeeze lever and valve assembly surmounting the tank and preferably provided with a lock pin and pressure gauge, a delivery hose connected to the valve assembly and terminating in a nozzle, and a cartridge holder connected to the hose upstream of the nozzle and containing a cartridge comprising a shaped, solid body having at least one exposed surface and comprising a solid, water soluble, coalesced mixture of water soluble, 40 fluoroaliphatic surfactant and compatible, water soluble, fluorine-free surfactant, which extinguisher,

upon activation, operates by discharging water from the tank through said cartridge holder to contact said surface to dissolve said surfactants and provide via the nozzle a foam of an aqueous film-forming foam ("AFFF") solution of relatively constant composition which can be applied to a body of flammable liquid to extinguish a fire thereof (viz., a "Class B" fire), or prevent such fire, by forming a tough,

45 durable, rapidly-forming and spreading aqueous film on the surface of the flammable liquid in a general manner described in the aforementioned patents. The extinguisher of this invention can also be used to more effectively combat a fire of solid combustible material, such as paper and wood (viz., a "Class A" fire), than a conventional hand portable 2.5-gallon water fire extinguisher. Such a conventional extinguisher can be readily converted, as described below, to an extinguisher of this invention for

50 combating both classes of fire.

The cartridges employed in the fire extinguishing systems of this invention can comprise a single, shaped body comprising the solid surfactant mixture. For example, in one embodiment of cartridge construction, the solid surfactant mixture is in the form of a sheet, with or without a reinforcing matrix (e.g., a sheet of needle felt), laid upon a non-woven, water-insoluble, water-permeable fabric and the 55 assembly rolled up as a coil and loaded as a cartridge in a sleeve, and the loaded sleeve inserted in a cartridge holder. More specifically, the solid surfactant mixture can be heated and pressed into a sheet, trimmed to size, placed in exact alignment on a piece of non-woven fabric, heated to soften, rolled up and inserted as a cartridge into a sleeve. In another example, the shaped body comprising the solid surfactant mixture is in the form of a single, solid cylinder with a star-shaped axial channel for water 60 passage. In another exbodiment, the cartridge is a water-permeable aggregation of a plurality (e.g. 50 to 20,000) of shaped bodies (comprising said solid mixture of surfactants) in the form of discrete pellets, beads, rods, etc., of relatively uniform size and shape, the water-permeability of the aggregation being due to the interstitial spaces between the plurality of shaped pellets, etc., providing channels for the water passed through the cartridge.

These types of cartridges, made up of one or more shaped bodies of the solid mixture of surfactants, produce aqueous film-forming solution of relatively constant composition as the surfactant mixture dissolves in the water stream supplied from the tank. The total surface area of the shaped body or bodies of surfactant mixture is a predetermined surface area which is sufficient to produce the AFFF solution with the requisite relatively constant composition at a given discharge rate. Said 5 predetermined surface area will vary, depending upon the particular solid surfactant mixture, the fabrication and formulation of the shaped body or bodies, the number, shape and size of the shaped body or bodies, and the volume, discharge rate, and temperature of the water in the tank. By "relatively constant composition", it is meant that the minimum concentration of the surfactants in the resulting solution during the discharge period of the fire extinguisher is at least about 50 percent, preferably at 10 least 55 percent, of the maximum concentration. To ensure said requisite composition, the amount of shaped body or bodies of the cartridge is such that there will generally be a residual amount of shaped body or bodies left in the cartridge holder after all the water is discharged from the tank. In the accompanying drawing: 15 Fig. 1 is a view in elevation of one embodiment of a fire extinguisher of this invention provided 15 with a loaded cartridge holder; Fig. 2 is a longitudinal view in partial cross-section of the loaded cartridge holder-nozzle of Fig. 1; Fig. 3 is an isometric view of one embodiment of a cartridge preform or layup which can be rolled up and loaded in the cartridge holder of Fig. 2; and 20 Fig. 4 is an isometric view of a cartridge assembled from the preform of Fig. 3. 20 Referring now to the drawing, and initially to Fig. 1 reference number 1 denotes one embodiment of a hand portable fire extinguisher of this invention comprising a tank 2 surmounted by an assembly comprising a squeeze lever 3, pressure gauge 4, valve (not visible), and lock pin 6, this assembly being connected to a delivery hose 7, the end of which is connected to a cartridge holder 8, which is 25 connected to an air aspirating nozzle 9. A conventional, Class A, portable, water fire extinguisher 25 (typically containing about 9.5 liters of water and about 2.8 liters of compressed gas, e.g. air or nitrogen) can be simply modified, for purposes of this invention, by removing its straight stream nozzle and replacing it with the combined cartridge holder-nozzle assembly shown in Figs. 1 and 2. Alternatively, a conventional Class A fire extinguisher can be modified by cutting its customary delivery 30 hose, interposing a cartridge holder of this invention, affixing the cartridge holder to the cut ends of the 30 hose with suitable clamps or the like, and replacing the customary nozzle with an air aspirating nozzle. Fig. 2 represents one embodiment of a loaded cartridge holder of this invention, wherein a split cylinder made of parts 10 and 11, threaded to engage one another, e.g. by means of a straight thread or spiral buttress thread such as described in NASA Tech Brief 71-10336 (September 1971), is 35 provided with an internal cartridge chamber 12 (e.g., about 250 cm³ in volume) adapted to contain 35 within sleeve 20 a cartridge 13 of the invention, the upstream end of cylinder part 11 having a fitting 5 adapted to receive the downstream end of hose 7 and the downstream end of cylinder part 10 being adapted for connection to an air aspirating nozzle 9, the upstream end of which is provided with air openings 15 and with a check valve 16 to prevent fluid, e.g., moist air or water, from flowing into the 40 cartridge holder via the nozzle. The exterior of the cartridge holder can be provided with flutes, as 40 shown, to enable the operator to firmly grasp the holder-nozzle unit during discharge of the extinguisher. In Fig. 3 there is illustrated a cartridge preform or layup 13' comprising a flat, rectangular porous substrate 21 on which is disposed a shaped body 22 of said fluoroaliphatic and fluorine-free 45 surfactants in the form of a slightly smaller, flat rectangular sheet being set back from one end and the 45 sides of the substrate 21, the other end of the sheet 22 being coincident with the other end of substrate 21. When the preform 13' of Fig. 3 is rolled upon itself it assumes the coil or roll form 13 shown in Fig. 4, the last loop of substrate 21 forming the exterior wall of the coil. Thus assembled, cartridge 13 can be inserted in a sleeve 20 (which can be considered as part of the cartridge), made of 50 plastic, metal, cardboard, phenolic-impregnated paper, etc., and placed in the cartridge chamber 12 of 50 cartridge holder 8 of Fig. 2, the porous substrate 21 serving both to separate the portions of the surface of shaped body 22 and to provide channels for the water to pass in contact with said surfaces as it flows from the tank to the nozzle. Optionally, as shown in Figs. 3 and 4, a second, short, porous substrate 23 can be placed on one end of the shaped body 22 to provide a central channel when the 55 preform 13' is rolled upon itself. 55 In operation, control of a fire is obtained by removing lock pin 6, squeezing operating lever 3 to open the valve and permit the pressurized water to flow via hose 7 into cartridge holder 8 and discharge from nozzle 9 a foam of an "AFFF" solution of relatively constant composition, viz., a foam of an aqueous film-forming foam solution generally containing 0.05 to 1 wt.% fluoro-aliphatic surfactant, 60 the weight ratio of fluoroaliphatic surfactant to fluorine-free surfactant being 10:1 to 1:25. The water, 60 in flowing through the cartridge holder, passes through the channels provided by the porous substrate 21, 23 in contact with the surface of the shaped body 22 to dissolve the mixture of surfactants. The surface area of the shaped body 22, and the rate of dissolution of each component, are relatively constant over the period of discharge. The 2.5 gallons (9.5 liters) of water will be discharged over a 45 65 to 90 second period at an initial pressure of about 7 kgf/cm2. The extinguisher can be recharged with

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about 9.5 liters of water and about 2.8 liters of compressed gas and with a new cartridge after drying the cartridge holder.

Where the cartridge used is a single, shaped body, the cartridge can be fabricated in a host of other forms so long as the surface area of the shaped body of surfactants exposed to the flow of water 5 from the tank does not significantly vary during dissolution of the surfactants. For example, the shaped body can be a solid cylinder with an axial channel having the shape of a star in transverse crosssection, the wall of the channel being exposed to the water discharged from the tank. In such embodiment, a porous substrate usually will not be required.

As described above, rather than loading the cartridge holder with a cartridge comprising a single, 10 shaped body of the solid surfactant mixture, the cartridge can comprise a water-permeable aggregation 10 of a plurality of shaped bodies, e.g., pellets, the surface area of which decreases during dissolution. Referring to Fig. 2, cartridge 13 can thus be replaced by an aggregation of a plurality of said shaped bodies. Sleeve 20 can be provided at least on its downstream end with a suitable, liquid-permeable or porous end-cap, such as one or more discs made of a low density, open, non-woven web, e.g., that 15 described in U.S. Pat. Nos. 2,958,593, 3,537,121 and that sold under the trademark "Scotch-Brite". Such discs can be fastened in place if desired, at their periphery to the sleeve, e.g., by a friction-fit or with a water-insoluble adhesive, e.g., a room temperature vulcanizable ("RTV") silicone rubber. Such end-caps will retain the aggregation in the sleeve during handling and during dissolution upon operation of the extinguisher.

The plurality of shaped bodies can be made by shaping the solid mixture of surfactants into the requisite size and shape by extrusion or pelletizing the solid mixture. In order to maintain channels for the water passing through a cartridge made of such shaped bodies, they are preferably fabricated with spherical or oblate shapes, i.e., bodies with mainly round or curved surfaces.

The shaped body or bodies of surfactants used in this invention, e.g. sheet 22 of Figs. 3 and 4, 25 comprises a solid, coalesced mixture of one or more water soluble fluoroaliphatic surfactants and one or more compatible, water soluble fluorine-free surfactants. The mixture is normally solid at ambient temperatures and does not become liquid below about 50°C.

The fluoroaliphatic surfactant contains one or more fluorinated aliphatic radicals (R_f) and one or more water solubilizing polar groups (Z) which are usually connected by suitable linking groups (Q).

Fluoroaliphatic surfactants especially useful are those disclosed in said U.S. Patent No. 3,562,156. The particular structure of the fluoroaliphatic surfactant is not critical; rather, it is the balance of the physical properties of the compound that determines its usefulness for the purpose. It is necessary that the combination of fluoroaliphatic radical and water solubilizing group be so balanced as to provide the surfactant with a solubility in water at 25°C. of at least 0.01 percent by weight. It is 35 preferred that the solubility in water be at least about 0.25 percent by weight. The surfactant must be sufficiently surface active to provide a surface tension of less than about 28 dynes/cm, preferably less than 23 dynes/cm, in aqueous solution at a concentration of about 0.25% or less.

If the fluoroallphatic surfactant is too soluble in hydrocarbon liquid, it will be extracted too rapidly from the aqueous film to provide sufficiently durable coverage. In general, this requires the presence of at least about 20 percent by weight of fluorine, i.e., carbon-bonded fluorine, in the surfactant. To possess these properties, the fluorinated aliphatic radical can be generally described as a fluorinated, saturated, monovalent, non-aromatic radical of at least 3 carbon atoms. The aliphatic chain may be straight, branched, or, if sufficiently large, cyclic and may include oxygen or trivalent nitrogen atoms bonded only to carbon atoms. A fully fluorinated radical is preferred, but hydrogen or chlorine atoms 45 may be present as substituents provided that not more than one atom of either is present for every two carbon atoms, and, preferably, the radical contains at least a terminal perfluoromethyl group. While radicals containing a larger number of carbon atoms will function adequately, compounds containing not more than about 20 carbon atoms are preferred since larger radicals usually represent a less efficient utilization of fluorine than is possible with shorter chains. Fluoroaliphatic radicals containing 50 about 5 to 12 carbon atoms are most preferred.

The water solubilizing polar group can be an anionic, a cationic, a non-ionic or ampholytic moiety or combinations thereof. Typical anionic groups would include CO₂H, CO₂M, SO₂M, SO₃H, SO₃M, OP(OH)2, and OP(OM)2, where M is a metallic ion, such as sodium, potassium, calcium, etc. Typical cationic groups would include NH2, NHR, where R is a lower alkyl group such as methyl, ethyl or butyl, 55 NR'3A', where R' is a lower alkyl group or hydrogen and A' is an anion, such as chloride, sulphate, phosphate, hydroxyl, etc. Typical non-ionic groups would include —NR₂→0 and those derived from polyethylene oxide and mixed polyethylene oxide-polypropylene oxide polyols. Typical mixed or ampholytic groups would include — $N(C_2H_4OH)_2$, — $NHC_2H_4NHC_2H_4NH_2$, — $N^+(CH_3)_2C_2H_4CO_2^-$, $[N^+(CH_3)_2C_2H_4COONa]OH^-$, ---N(CH₃) $(C_2H_4CO_2H)\rightarrow 0$, and the like.

The linking group is a multivalent, generally divalent, linking group such as alkylene, arylene, sulfonamidoalkylene, carbonamidoalkylene, and the like. In some instances more than oen fluoroaliphatic radical may attach to a single linking group and in other instances a single fluoroaliphatic radical may be linked to more than one linking group or may be linked by a single linking group to more than one polar solubilizing group.

A particularly useful class of fluoroaliphatic surfactants which can be used in this invention are

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Representativ fluoroaliphatic surfactants useful in this invention include:

5	C ₈ F ₁₇ SO ₃ K C ₆ F ₁₃ SO ₂ N(CH ₂ CHOHCH ₂ SO ₃)C ₃ H ₆ N ⁺ (CH ₃) ₂ C ₂ H ₄ OH C ₈ F ₁₇ SO ₂ NHCH ₂ C ₆ H ₄ SO ₃ Na C ₈ F ₁₇ SO ₂ NHC ₆ H ₄ SO ₃ Na	5
10	$C_{6}^{\circ}F_{13}^{\circ}SO_{2}^{\circ}N(C_{3}H_{6}SO_{3}^{\circ})C_{3}H_{6}N^{+}(CH_{3})_{2}C_{2}H_{4}OH$ $C_{7}F_{15}CONHC_{3}H_{6}N^{+}(CH_{3})_{2}C_{2}H_{4}COO^{-}$ $C_{8}F_{17}C_{2}H_{4}SC_{2}H_{4}CONHC(CH_{3})_{2}CH_{2}SO_{3}Na$ $C_{8}F_{17}SO_{2}N(C_{2}H_{5})C_{2}H_{4}P(O)(OH)_{2}$ $C_{6}F_{13}SO_{2}NHC_{3}H_{6}N^{+}(CH_{3})_{3}CI^{-}$	10
15	$C_8F_{17}SO_2NHC_3H_6N^+(CH_3)_3O_3SOC_2H_5$ $(CF_3)_2CF(CF_2)_6COOH_1NC_2H_5$ $C_7F_{15}COOH_1NC_3H_6N^+(CH_3)_2C_2H_4COO^ C_7F_{15}CONHC_3H_6N(CH_3)_2\rightarrow O$ $C_8F_{17}SO_2N(C_2H_5)CH_2CO_4K$	15
20	C ₈ F ₁₃ C ₂ H ₄ SO ₂ N(CH ₃)C ₂ H ₄ N ⁺ (CH ₃) ₂ C ₂ H ₄ COO C ₆ F ₁₃ SO ₂ N(CH ₂ CHOHCH ₂ SO ₃ Na)C ₃ H ₆ N(CH ₃) ₂ C ₈ F ₁₇ C ₂ H ₄ SCH(CH ₂ COONa)COONa C ₈ F ₁₇ C ₂ H ₄ SC ₂ H ₄ CONHC ₂ H ₄ N ⁺ (CH ₃) ₃ Cl C ₁₀ F ₂₀ HOC ₆ H ₄ SO ₃ Na	20
25	$(CF_3)_2^*CF(CF_2)_4^*CONHC_2H_4SO_3Na$ $[C_6F_{13}SO_2NHC_2H_6N^+(CH_3)_2C_2H_4OH]OH^-$ $[C_6F_{13}SO_2N(CH_2CH_2OH)C_3H_6N^+(CH_3)C_2H_4OH]OH^-$ $C_6F_{13}SO_2N(CH_2CH_2OH)C_3H_6N(CH_3)_2$	25

and mixtures thereof.

The water soluble, fluorine-free surfactants used in this invention are those which are synthetic, imputrescible, hydrocarbon-congruous organic compounds which are water soluble to at least about 0.02 percent by weight in water at 25°C. and are capable of promoting the film-forming ability of a normally non-film-forming, aqueous fluorocarbon surfactant solution. Such surfactants substantially completely emulsify at least one phase of a mixture of equal volumes of cyclohexane and water at a concentration of about 0.1 to 10 wt.% of the water. Additionally, the fluorine-free surfactants used in this invention must be compatible with the fluoroaliphatic surfactants. Compatibility here means that the two types of surfactants do not interact to produce an inactive product.

The fluorine-free surfactants particularly useful in this invention are those described in the aforementioned patents and they can be selected on the basis of the tests described in U.S. Pat. No. 3,772,195. Representative fluorine-free surfactants useful in the practice of this invention include:

$$C_8H_{17}OSO_3Na$$

 $C_{10}H_{21}OSO_3Na$
 $C_{12}H_{25}OSO_3Na$
 $C_{10}H_{21}SO_3K$

$$\begin{array}{c} \mathsf{C}_{12}\mathsf{H}_{25}\mathsf{N}(\mathsf{CH}_2\mathsf{COON8})_2\\ \mathsf{C}_8\mathsf{H}_{17}\mathsf{C}_6\mathsf{H}_4\mathsf{O}(\mathsf{C}_2\mathsf{H}_4\mathsf{O})_{30}\mathsf{H} \\ \mathsf{C}_{12}\mathsf{H}_{25}\mathsf{N}^+(\mathsf{CH}_3)_2\mathsf{C}_2\mathsf{H}_4\mathsf{SO}_3^-\\ \mathsf{C}_{12}\mathsf{H}_{25}\mathsf{N}^+(\mathsf{CH}_3)_2\mathsf{CI}^-\\ \mathsf{C}_{12}\mathsf{H}_{26}\mathsf{N}^+(\mathsf{CH}_3)_3\mathsf{CI}^-\\ \mathsf{(C}_8\mathsf{H}_{17}\mathsf{O})_2\mathsf{PO}_2\mathsf{Na} \\ \mathsf{HO}(\mathsf{C}_2\mathsf{H}_4\mathsf{O})_a(\mathsf{C}_3\mathsf{H}_6\mathsf{O})_b(\mathsf{C}_2\mathsf{H}_4\mathsf{O})_c\mathsf{H}, \, \mathsf{MW} \, 6500\\ \mathsf{C}_{12}\mathsf{H}_{26}\mathsf{O}(\mathsf{C}_2\mathsf{H}_4\mathsf{O})_4\mathsf{C}_2\mathsf{H}_4\mathsf{OSO}_3^-\mathsf{NH}_4^+\\ \mathsf{C}_8\mathsf{H}_{17}\mathsf{SC}_2\mathsf{H}_4\mathsf{CONHC}(\mathsf{CH}_3)_2\mathsf{CH}_2\mathsf{SO}_3\mathsf{Na}\\ \mathsf{C}_{12}\mathsf{H}_{25}\mathsf{SO}_2\mathsf{N}(\mathsf{CH}_2\mathsf{COO}^-)\mathsf{C}_3\mathsf{H}_6\mathsf{N}^+(\mathsf{CH}_3)_3\\ \mathsf{C}_{12}\mathsf{H}_{26}\mathsf{N}(\mathsf{CH}_3)_2\to\mathsf{O} \end{array}$$

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and mixtures thereof. Certain fluorine-free silicone surfactants are known to be useful in forming AFFF solutions and they can be used here too.

In general, the weight ratio of fluorine-free surfactant to fluoroaliphatic surfactant is in the range of 1:25 to 10:1.

The formulation of the shaped body or bodles can contain, in addition to the two types of surfactants, various adjuvants which aid in the processing or formulation of the shaped body (e.g., solid polyethylene glycols, or methoxy polyethylene glycols, with number average molecular weights of 1000 to 20,000, preferably 1000 to 6000), foam stabilizers (e.g., polysaccharide foam stabilizers) which stabilize the foam when applied to lower alcohols, ketones, and other flammable polar liquid, 10 adjuvants which modify the softening temperature of the shaped body (e.g., sorbitol), effervescents which aid dissolution) e.g., citric acid with sodium blcarbonate), adjuvants commonly used in preparing AFFF solutions (provided such adjuvants are compatible with the particular surfactant combination used and do not unduly lower the softening point of the desired shaped body below 50°C), and antioxidants and biocides, such as fungicides, which enhance stability and shelf life of the shaped body 15 or bodies. In some cases individual fluoroaliphatic surfactants and fluorine-free surfactants can provide the desired AFFF solutions, but more frequently mixtures of two or more of each type of the surfactants are more readily available and can be used to provide more desirable AFFF solutions.

A particularly useful formulation for the shaped body comprises the following:

20	C ₆ F ₁₃ SO ₂ N(CH ₂ CHOHCH ₂ SO ₃)CH ₂ CH ₂ CH ₂ CH ₂ N+(CH ₃) ₂ CH ₂ CH ₂ OH	10—20 wt.% 5—20	20
20	C ₈ F ₁₇ SO ₃ K C ₁₀ H ₂₇ OSO ₃ Na	40—80 0—20	
	HO(CH ₂ CH ₂ O) _m H (ave. mol. wt 3000 to 4000) HOCH ₂ (CHOH) ₄ CH ₂ OH (sorbitol)	0—20	

Solid mixtures of surfactants can be prepared by spraying an aqueous solution containing the 25 fluoroaliphatic surfactant and fluorine-free surfactant (in the ratios described above) in a spray drier, such as that manufactured by the Niro Atomizer, Inc. and sold under the trademark "Niro", this spray drier having a 1.26 meter diameter. The spray drier can be operated with an air flow at about 7.5 m³/min. using an air inlet temperature of about 80°C and an exit air temperature of about 40°C. The solution can be metered onto a high speed (e.g. 24,000 RPM), rotating slotted disc at the top of the unit, which atomizes the solution into tiny droplets. The droplets are dehydrated by the flowing air and the resulting solid particles are collected by means of an air cyclone separator. The particles can be formed into the requisite shaped body or bodies by various means, such as by extrusion, calendering, molding, and the like.

When the cartridge is a single, shaped body, the shaped body is preferably in the form of a sheet (which can be corrugated, embossed, etc., on one or both surfaces to increase surface area) which is rolled up with a water insoluble, water permeable, open, resilient, three-dimensional web, such as shown in the drawing. A particularly useful substrate for this purpose is the low-density, open, nonwoven, three-dimensional web formed of many interlaced, randomly disposed, flexible, durable, tough, organic fibers which are firmly bonded together at points where they intersect and contact one another by globules of an organic binder, such fibrous material being described in U.S. Pat. Nos. 2,958,593 (Hoover) and 3,537,121 (McAvoy). Commercial articles of such fibrous material with 70—95% void volume and made of thermoplastic fibers are sold under the trademark "Scotch-Brite", e.g., "Scotch-Brite" Type A made of nylon 66 having a 12-15 denier (12-15 g/9000 meters).

Alternatively, the single, shaped body can be prepared by saturating a water permeable, water 45 insoluble, reinforcing matrix, e.g., a porous, fibrous web, such as felt, wool batting, etc., with a solution of the mixture of surfactants and removing the solvent. The impregnated web or matrix can then be rolled up with a porous substrate, such as the aforementioned "Scotch-Brite".

Objects and advantages of this invention are illustrated in the following examples in which the amounts given are parts by weight and the water is deionized water, unless indicated otherwise. The 50 controls used were tap water solutions of the particular formulations described, the concentrations of such solutions being indicated in the tables. Where film speeds are reported, they were obtained by placing 2 drops of the AFFF solution onto the surface of cyclohexane contained at room temperature (about 22°C) in a 5-cm. dlameter petri dish and measuring the time for the film to cover the surface.

Example 1

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The following ingredients were combined, stirred, and heated (85°C.) for about 30 minutes to form a homogeneous solution.

Table !

60	2.95	$C_6F_{13}SO_2N(CH_2CHOHCH_2SO_3^-)CH_2CH_2CH_2CH_2N^+(CH_3)_2CH_2CH_2OH^6$ $C_8F_{17}SO_3K$	60
	0.5	C ₁₂ H ₂₅ OSO ₃ Na	

Table I (contd.) PART IN

parts	
11.4	C ₁₀ H ₂₁ OSO ₃ Na
22.1	HÖ(CH,CH,O),Hb.
22.1	CH ₃ O(CH ₂ CH ₂ Ö) _n H°
35.0	Water
	11.4 22.1 22.1

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- Prepared, following the procedure of Example 7 of Australian Patent Specification 38028/72. by reaction of C₈F₁₃SO₂N(Na)(CH₂)₃N⁺(CH₃)₂CH₂CH₂OH with sodium-2-hydroxy-3-chloropropanesulfonate instead of propanesultone.
- "Carbowax" polyethylene glycol 4000 (ave. molecular weight about 3000-3700). 10

"Carbowax" methoxy polyethylene glycol 2000 (ave. molecular weight about 1900).

The hot solution (211 g) was used to saturate a 24 cm x 14 cm piece of needle felt (65/35 polyester/viscose rayon, density 0.088 g/cm³, 0.32 cm thick). The resulting impregnated felt was dried at about 100°C, for 6 hrs. to remove most of the water. The dried felt containing the coalesced mixture 15 was trimmed to about 22 cm x 14 cm and placed on a 15 cm x 24 cm x 0.5 cm piece of "Scotch-Brite" Type A fabric and the combined layers rolled up tightly, with the "Scotch-Brite" fabric on the outside, to form a cartridge 15 cm long with a diameter of about 4 cm. The cartridge was fitted snugly into a galvanized steel pipe (4 cm x 15 cm long) threaded on both ends. Pipe caps with adapters for hose were placed on each end of the pipe. The loaded pipe, or cartridge holder, was inserted in the hose line 20 of a standard hand portable, 2.5-gallon, water fire extinguisher, the nozzle of which was replaced with an air-aspirating foam nozzle having a flow rate of about 20 liters/min. at 7 kgf/cm2. The extinguisher was filled with about 9.5 liters of tap water, pressurized to about 7 kgf/cm2. The extinguisher was discharged to yield an AFFF solution having, as shown below, nearly uniform concentration of solute over the discharge period as determined from refractive index measurements of samples taken at 10

Table II Time, solute conc., Film speed, g/Isec. sec. 10 9 30 20 8 30 8 30 8 40 18 8 50 7 60 35 7 70 35 80 7 24

Example 2

25 sec. intervals.

The following Ingredients were combined, stirred and heated (85°C) for about 30 minutes to form a homogeneous solution.

40		Table III	40
45	Parts 8.6 4.3 25.7 0.8 12.8 12.8 35.0	$C_6F_{13}SO_2N(CH_2CHOHCH_2SO_3^-)CH_2CH_2CH_2N^+(CH_3)_2CH_2CH_2OH C_8F_{17}SO_3K C_{10}H_{21}OSO_3Na C_{12}H_{25}OSO_3Na HO(CH_2CH_2O)_nH^a$ HO(CH_2CH_2O)_nH^a water	45
50	^{a.} "Cal ^{b.} Sort	rbowax" 4000 pitol	50

The aqueous solution (1600 g) was spray dried in a "Niro" utility spray drier using the abovedescribed conditions and 1 wt% fumed silica ("Cabosil" MS-7) was added to yield a free-flowing powder (8 to 65 micrometers).

A 150 g sample of the powdered product was pressed in a platen press at 70 kgf/cm² to form a 55 coalesced, solid flat sheet having an average thickness f about 0.34 cm. This sheet was trimmed to the dimensions of about 13 cm x 19 cm and found to weigh 118 g. It was placed on a 15 cm x 24 cm piece of "Scotch-Brite" Type A web centered in the narrow dim insion, and positioned with one end (A) coincident with the end of the w b, and with a second piece 2.5 cm x 15 cm of the web placed over the

solid sheet at end A so as to make a sandwich construction. After warming this construction for 1 hour at 66°C. to soften the solid sheet, it was rolled up tightly from end A (with the larger piece of web on the outside) and inserted into a 4 cmx 15 cm steel pipe. The pipe was connected to a 2.5-gallon water fire extinguisher as described in Example 1. The extinguisher was filled with about 9.5 liters of tap water, pressured to about 7 kgf/cm² with compressed nitrogen gas, and discharged completely over a period of 68 seconds. An AFFF solution of good quality and fairly uniform concentration was produced as shown by the following data obtained on samples collected during discharge.

Table IV

				Film	
10	Time,	Refractive	Solute conc.,	speed,	10
	sec.	index n ²⁰	g/I	SBC	
	2	1.3338	7	10	
	10	1.3338	7	9	
	20	1.3337	6	11	
15	30	1.3337	6	10	15
. •	40	1.3337	6	15	
	50	1.3337	6	15	
	60	1.3337	6	23	
	68	1.3339	8	6	
20	Tap Water	1.3330	0		20
	Control	1.3339	8	6	

The cartridge was taken apart and the web unrolled. The remaining solid formulation weighed 55 g, showing that 63 g of solid formulation had dissolved, apparently uniformly, resulting in an average concentration of 6.6 g/liter.

25 Example 3

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The following ingredients were combined, stirred and heated at about 80°C for about 30 min. to form a homogeneous solution.

Table V

	Parts		00
30	88.4	C ₆ F ₁₃ SO ₂ N(CH ₂ CHOHCH ₂ SO ₃ ⁻)CH ₂ CH ₂ CH ₂ CH ₂ N ⁺ (CH ₃) ₂ CH ₂ CH ₂ OH	30
	44.2	C _R F ₁₇ SO ₃ K	
	172	C ₁₀ H ₂₁ OŠO ₃ Na	
	7.5	C ₁₂ H ₂₅ OSO ₃ Na	
	331.5	HÖ(CH,CH,O),H³·	
35	331.5	CH ₃ O(CH ₂ CH ₂ Ö) _n H ^{b.}	35
	524.9	Water	

- a. "Carbowax" 4000
- b. "Carbowax" 2000

About 210 g of the above formulation (a 65% solution) was poured over a 24 cm x 14 cm piece of needle felt (same as that of Example 1) in a 15 cm x 25 cm glass tray. The tray was placed in a 110°C. forced air oven and the water allowed to evaporate from the composition over a 6.5 hour period. the solid composite was turned over and heated in a vacuum oven at 50°C. for 2 hours, trimmed to 22 cm x 14 cm, placed on a 24 cm x 15 cm plece of "Scotch-Brite" Type A and the combined layers rolled up tightly, with the web on the outside, and inserted into a 15 cm piece of 4 cm threaded steel pipe.

The pipe was connected to a hose line of a 2.5-gallon water fire extinguisher as described in Example 2, the extinguisher filled with about 9.5 liters of tap water, and pressurized to about 7 kgf/cm² with nitrogen gas.

An approximately 2.5 cm layer of heptane (over a layer of water) in a 4.65 m² square steel pan was ignited and allowed to burn for 30 sec., then extinguished in a period of 71 seconds using essentially all of the contents of the fire extinguisher described above. Details of events are given below.

Tabl VI

	Time, min:sec	Event	
55	0:00	ignition	55
	0:30	begin extinguishing fire	
	0:40	15% of fire was extinguished	
	0:50	40% ,, ,,	

		Tab	le VI (c nto	d.)		
	Time,	•	_			
	min:sec	0.00/	Event			
_	0:60	90%	11	"		
5	1:10	97%	"	"		
	1:20	99%	"	"		
	1:30	98%	11	"		
	1:40	99%	"	**		
	1:41	100%	"	"		
)	10:56	attempte	d to reignite	e heptane		
		with to	orch; a mino	or transient		
		flame v	was observe	ed, but a fire		
		could r	ot be susta	iined.		
5	Weighing of the pipe content in the felt carrier, and that 110 g		• •		18 g of solid material remained	
	Example 4					
	A solution was prepared fro	m the following	g ingredient	s.		
		· т	able VII			
)	Parts 5.0 C ₆ F ₁₃ SO ₂ N(CH ₂ CH0	OHCH,SO, ⁻)CH	,CH,CH,N+	(CH ₃) ₂ CH ₂ CH	,ОН	
	2.5 C ₈ F ₁₇ SO ₃ K					
	9.75 Mixture ^{6,} of 42 part 0.4 C ₁₂ H ₂₅ OSO ₃ Na	18 C ₈ H ₁₇ USU ₃ N	a anu o7 pa		J ₃ IVa	
	24.5 HÖ(CH ₂ CH ₂ O) ₀ Hb·					
j	24.5 CH ₃ O(CH ₂ CH ₂ O) _n H					
	33.35 water	•				
	33.35 Water					
	B. This mixture sold under	the trademark	"Stepan" 6	70-15		
	 This mixture sold under "Carbowax" 4000 	the trademark	"Stepan" 6	370-15		
	^{8.} This mixture sold under	the trademark	"Stepan" 6	370-15		
	 This mixture sold under "Carbowax" 4000 "Carbowax" 2000 					
כ	 This mixture sold under "Carbowax" 4000 "Carbowax" 2000 The pH of the solution was 	adjusted to 9.0	with 10% a	aqueous sodic	ım hydroxide. The foam and	
ס	B. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast	adjusted to 9.0 ured in accorda	with 10% a	aqueous sodio	on F-24385B Amendment 1,	
כ	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution con-	adjusted to 9.0 ured in accorda	with 10% a	aqueous sodio	on F-24385B Amendment 1,	
ס	B. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast	adjusted to 9.0 ured in accorda taining 100 g o	with 10% a nce with MI f solids in 9	aqueous sodio	on F-24385B Amendment 1,	
	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution con-	adjusted to 9.0 ured in accorda taining 100 g o	with 10% a	aqueous sodio	on F-24385B Amendment 1, iter, were found to be as	
	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows.	adjusted to 9.0 ured in accorda taining 100 g o T	with 10% a nce with MI f solids in 9	aqueous sodio	on F-24385B Amendment 1, iter, were found to be as Interfacial tension	
	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows.	adjusted to 9.0 ured in accordant taining 100 g o T	with 10% a nce with MI f solids in 9 able VIII	aqueous sodic IL Specification.5 liters of wa	Interfacial tension between cyclohexane	
	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows.	adjusted to 9.0 ured in accordant taining 100 g o T 25% drain time ^(b) ,	with 10% a nce with MI f solids in 9 able VIII Surface	aqueous sodication.5 liters of water tension,	In F-24385B Amendment 1, iter, were found to be as Interfacial tension between cyclohexane and water	
	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows.	adjusted to 9.0 ured in accordantaining 100 g o T 25% drain time ^(b) , min.	with 10% a nce with MI f solids in 9 able VIII Surface dynes/cm	aqueous sodication.5 liters of watension, at 22°C	Interfacial tension between cyclohexane and water dynes/cm at 22°C	
	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows.	adjusted to 9.0 ured in accordant taining 100 g o T 25% drain time ^(b) ,	with 10% a nce with MI f solids in 9 able VIII Surface dynes/cm	aqueous sodication.5 liters of water tension,	In F-24385B Amendment 1, iter, were found to be as Interfacial tension between cyclohexane and water	
5	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows. Foam expansion (a) 8.3 (a) Ratio of foam volume to the solution of the solution confollows.	adjusted to 9.0 ured in accordantaining 100 g o T 25% drain time ^(b) , min. 4.8	with 10% a nce with MI f solids in 9 able VIII Surface dynes/cm 15 me.	aqueous sodication.5 liters of watension, at 22°C	Interfacial tension between cyclohexane and water dynes/cm at 22°C	
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5	a. This mixture sold under b. "Carbowax" 4000 c. "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows. Foam expansion (a) 8.3 (a) Ratio of foam volume to (b) Time for 25% of liquid	adjusted to 9.0 ared in accordant taining 100 g of the state of the st	with 10% a nce with MI f solids in 9 able VIII Surface dynes/cm 15 me. ulk foam.	aqueous sodication.5 liters of wastension, at 22°C 5.8	Interfacial tension between cyclohexane and water dynes/cm at 22°C 4.2	
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5	This mixture sold under "Carbowax" 4000 "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows. Foam expansion (a) 8.3 (a) Ratio of foam volume to Time for 25% of liquid The above formulation was gave 0.39 g of coalesced solids pages.	adjusted to 9.0 ared in accordant taining 100 g of tainin	with 10% a nce with MI f solids in 9 able VIII Surface dynes/cm 15 me. ulk foam. dle felt (sam ying. A 10 c	aqueous sodication.5 liters of wastension, at 22°C s.8	Interfacial tension between cyclohexane and water dynes/cm at 22°C 4.2 Example 1) at a level which ece of the dried impregnated	
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5 0 5	This mixture sold under "Carbowax" 4000 "Carbowax" 2000 The pH of the solution was surface activity properties, meast May 16, 1969, on a solution confollows. Foam expansion (a) 8.3 All Ratio of foam volume to the solution was gave 0.39 g of coalesced solids properties was placed on a 10 cmx2 dcmx10 cm long pipe cartridge gallon water fire extinguisher, the Example 1. Samples were collect properties determined to be as for the solution of the solution was gave 0.39 g of coalesced solids properties determined to be as for the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution of the solution water fire extinguisher, the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution of the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution water fire extinguisher, the example 1. Samples were collect properties determined to be as for the solution water fire extinguisher.	adjusted to 9.0 Ired in accordate taining 100 g of taini	with 10% ance with MI foolids in 9 able VIII Surface dynes/cm 15 able felt (same ving. A 10 concentrate in the lead, pressurange and concentrate in the lead of the lea	tension, at 22°C 3.8 Type A we cartridge horized and disconcentration of the conc., g// 10 9	Interfacial tension between cyclohexane and water dynes/cm at 22°C 4.2 Example 1) at a level which ece of the dried impregnated b and rolled up and placed in a lder was attached to a 2.5- charged as described in of solute and film-forming Film speed	

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A fire test with 9.5 liters of the control solution above, discharged over a 65 second period, from a 2.5-gallon fire extinguisher having an air aspirating nozzle, gave control of a test fire in 35 seconds and total extinguishment in 63 seconds. The test fire was a 4.6 m² heptane fire run by the method described in Underwriters Laboratories Standard 711.

5	Example 5 The following ingredients were combined, stirred and heated at about 50°C. to form a	!
	homogeneous solution with a pH of 4.2.	

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	e Ji		

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	Parts		·
10	42.1	C ₆ F ₁₃ SO ₂ NHC ₃ H ₆ N ⁺ (CH ₃) ₃ CI ⁻	10
	21.05	$C_8F_{17}SO_2NHC_3H_6N^+(CH_3)_3CI^-$	
	8.3	C ₇ F ₁₅ COÖ ⁻ H ₃ N ⁺ C ₃ H ₆ N ⁺ (CH ₃) ₂ C ₂ H ₄ COO ⁻	
	145.8	HO(C ₂ H ₄ O) _a (C ₃ H ₆ O) _b (C ₂ H ₄ O) _c H ^{6.}	
	70.9	HO(CH ₂ CH ₂ O) _n H ^{b.}	
15	70.9	HOCH₂(CHOH)₄CH₂OH°·	15
	5.9	CH ₃ COOH	
	3.5	CH ₃ COONa	
	530.5	water	
	a. "PL	.URONIC" F-77, molecular weight 6500	
20	ь. "Ca	ırbowax'' 4000	20
		bitol	

This solution (366.2 g) was used to saturate a piece of needle felt fabric (same as used in Example 1) in a glass tray. The dimensions, drying procedures, and cartridge preparation were the same as described in Example 3. The weight of solid, coalesced surfactant mixture in the cartridge was 117.5 g or 0.42 g/cm² of the impregnated felt.

The cartridge was attached to a 2.5 gallon water fire extinguisher, filled, pressured and discharged as described in Example 1. An AFFF solution was produced having an effective and nearly uniform concentration of solute over the discharge period as determined from the solute concentration in complex taken at about 10 sec. intervals

	in samples taken at about 10 sec. intervals.			30
30		Table XI	_	30
•	Time,	Refractive	Solute conc.,	
	sec.	index n _D ²⁰	g/I	
	2	1.3333	3	
	10	1.3333	3	
35	20	1.3333	3	35
00	30	1.3333	3	
	40	1.3333	3	
	50	1.3333	3	
	60	1.3333	3	•
40	70	1.3333	3	40
40	76	1.3335	4	
	Tap Water	1.3330	_	
	Control	1.3339	8	

Foam expansion and 25% drain time were measured between the 2 and 10 sec. intervals and found to be 2.4 and 3.5 minutes, respectively. Analysis of the cartridge contents after the test showed that 25.2 g of the 117.5 g of solid material had been utilized during discharge. Calculations indicate that 2.7 g solute per liter should have been present in the discharged solution, which is very close to the value obtained. While the AFFF solution was quite dilute, it was adequate to retard vaporization of volatile solvents.

50 Example 6

The following ingredients were combined using agitation and warming.

Table XII

55	Parts 11.2 22.4 350.8	$C_8F_{17}SO_3K$ $C_8F_{13}SO_2N(CH_2CHOHCH_2SO_3^-)CH_2CH_2CH_2N^+(CH_3)_2CH_2CH_2OH$ Sodium alkyl sulfates ("Polystep" B-25, a 38.6% aqueous solution of $C_8H_{17}OSO_3Na$, $C_{10}H_{21}OSO_3Na$, and $C_{12}H_{25}OSO_3Na$ in a weight ratio f about 2:75:23)	55
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Water was evaporated to yield 305.3 g of solution. About 270 g was poured into a 15 cm x 25 cm glass tray containing a 24 cm x 14 cm piece of needle felt (same as used in Example 1). The tray and

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contents wer heated at 108°C. for 13 hrs. to remove essentially all of the remaining water (within 2 g of the expected dry weight, i.e. about 1.5% water content). The dried felt was pressed in a platen press at 25°C., trimmed to 13 cm x 19 cm, placed on a 15 cm x 24 cm piece of non-woven fabric, warmed to soften the coalesced solids, and the combined materials rolled up and inserted int 4 cm x 15 cm threaded steel pipe and fitted with hose adapters.

The resulting cartridge was connected to a standard 2.5-gallon, water fire extinguisher, the extinguisher filled with about 9.5 liters tap water, pressured to 7 kgf/cm² and discharged completely over a period of 64 sec. An effective AFFF solution of fairly uniform composition was produced as shown by the following data obtained on samples collected during discharge.

10			Table	XIII			10
٠	•		:		Surface	Interfacial tension between cyclohexane	
15			Solute	FIlm	tension,	and water,	15
	Time,	Refractive	conc.,	speed,	dynes/cm	dynes/cm	
•	sec.	Index n ²⁰	<i>g</i> //	sec.	at 22°C	at 22°C	
	2 .	1.3341	10	60(a)	17.6	4.0	
	10	1.3337	6	60(b)			
20	20	1.3336	5.	21	18.0	3.3	20
	30	1.3337	6	45		•	
	40	1.3336	5	14	18.0	3.3	
	50	1.3336	5	18			
	60	1.3336	5	55	18.0	3.2	
25	64 ·	1.3337	6	35			25
	Control	1.3339	8	8	17.6	4.2	
	Tap Water	1.3330	0		· —		

- (a) 25% of cyclohexane surface covered in 60 sec.
- (b) 90% of cyclohexane surface covered in 60 sec.

30 Example 7

The following solid ingredients were sieved individually through a screen and mixed together.

- 11.3 g $C_0F_{17}SO_3K$
- 22.7 C₆F₁₃SO₂N(CH₂CHOHCH₂SO₃)CH₂CH₂CH₂CH₂CH₂CH₂OH
- 135.0 Sodium alkyl sulfates ("Polystep" B-25 dried solids)
- 35 The mixture was warmed to 100°C. and pressed several times on a platen press to yield a solid, pale amber, waxy coalesced sheet having an average thickness of 0.34 cm. After trimming, the solid sheet had the dimensions 13 cm x 19 cm and weighed 114.5 g. It was warmed to 110°C. to soften, placed on a 15 cm x 24 cm piece of "Scotch-Brite" Type A web and the combined layers rolled up and inserted in the pipe assembly described earlier. The finished cartridge was connected to a standard 2.5 gallon
- water fire extinguisher, the extinguisher filled with 9.5 liters tap water, pressurized to 7 kgf/cm² and discharged completely over a period of 57 seconds. An effective AFFF solution of quite uniform composition was produced as shown by the following data obtained on samples collected during discharge.

aloute go.		Table	XIV			
	-			Surface	Interfacial tension between cyclobexane	45
		Solute	Film		and water,	
Time,	Refractive	conc.,	speed,	dynes/cm	dynes/cm	50
sec.	index n ₀ 20	g/I	sec.	at 22°C	at 22°C	
2	1.3338	8	11			
10	1.3338	8	17	16.4	3.5	
20	1.3338	8	16			
30	1.3337	7	12	16.5	3.4	55
40	1.3337	7	10		_	
50	1.3337	7	9	16.7	3.4	
57	1.3339	8	11			
Control	1.3339	8	5	16.2	3.9	
Tap Water	1.3330	0		_	_	60
	sec. 2 10 20 30 40 50 57 Control	Time, Refractive sec. index n _D ²⁰ 2 1.3338 10 1.3338 20 1.3338 30 1.3337 40 1.3337 50 1.3337 57 1.3339 Control 1.3339	Table Time, sec. index n _D ²⁰ g/l 2 1.3338 8 10 1.3338 8 20 1.3338 8 30 1.3337 7 40 1.3337 7 50 1.3337 7 57 1.3339 8 Control 1.3339 8	Table XIV Time, sec. index n ₀ ²⁰ g/l sec. 2 1.3338 8 11 10 1.3338 8 17 20 1.3338 8 16 30 1.3337 7 12 40 1.3337 7 10 50 1.3337 7 9 57 1.3339 8 11 Control 1.3339 8 5	Table XIV Surface Solute Film tension, Time, Refractive conc., speed, dynes/cm sec. index n ₀ ²⁰ g/l sec. at 22°C 2 1.3338 8 11 — 10 1.3338 8 17 16.4 20 1.3338 8 16 — 30 1.3337 7 12 16.5 40 1.3337 7 10 — 50 1.3337 7 9 16.7 57 1.3339 8 11 — Control 1.3339 8 5 16.2	Table XIV

The weight of solid sheet of coalesced solids remaining after discharge was 53.3 g., indicating that 61.2 g had dissolved corresponding to an average concentration of 6.5 g f solute per liter.

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Example 8

One hundred ninety grams of the powdered surfactant product described in Example 2 were combined and mixed with 19 g of a powdered polysaccharide gum (K 8A13 made by Kelco Division of Merck & Co.) and the resulting mixture pressed in a platen press at 70 kgf/cm² to form a coalesced, 5 solid, flat sheet. Two sheets 0.25 cm thick were prepared and trimmed to the dimensions 13.3 cm x 14.0 cm (wt. 128 g). Using a mold with a saw-tooth pattern (0.18 cm deep valleys, 0.35 cm. between peaks), grooves were pressed into both surfaces of the sheets at right angles to the long dimension. These sheets were placed end-to-end, with the short dimension abutting, on a piece of "Scotch-Brite" fabric, warmed to soften, and the construction rolled up at right angles to the direction 10 of the grooves in the sheets to form a cartridge as described in Example 2. The cartridge was inserted into a polyvinyl-chloride plastic tube having the dimensions: 15.2 cm in length, 4.48 cm outside diameter, 4.25 cm inside diameter. This assembly was placed in an acrylonitrile/butadiene/styrene (ABS) plastic cartridge holder-nozzle assembly similar to that shown in Fig. 2 (the cartridge holder had a cavity 15.2 cm long and an inner diameter of 4.50 cm). This assembly was connected to a 2.5-gallon, 15 hand portable water fire extinguisher, the extinguisher filled with about 9.5 liters of tap water at 21°C, pressured to about 7 kgf/cm² with nitrogen, and discharged completely over a period of 66.5 sec.

Foam samples were taken at the discharge intervals shown and several properties of these samples measured and are summarized below.

				Table	XV				20
20	Time,	Refractive	Solute conc.,	Foam expan-	25% drain time,	Surface tension, dynes/cm	Interfacial tension, dynes/cm 22°C	Film speed, sec.	20
·25	<i>sec.</i> 5 40	5 1.337 40 1.33355	6.1 4.8	sion 8.6 7.8	min. 5.3 2.2	22°C 16.8 16.8 16.7	3.3 3.3	8 13 3.5	25
	Control Tap Water	1.3338 1.3330	7.0				-	_	

The weight of the residual solid, coalesced sheets in the cartridge holder weighed 71 g, showing that 57 g had dissolved (44%).

In another example, which was like that described above (except that the polysaccharide was dried to remove adventitious moisture, and ratio of the powdered surfactant product/dried polysaccaride was 188 g/12 g), a higher percentage of the solid formulation dissolved: 74 g out of 128 g. (58%).

Example 9

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The following ingredients were combined stirred and heated (about 85°C.) for about 30 min. to 35 form a homogeneous solution.

- "Richonol" 7227
- "Niaproof' Anionic 7 50
 - "Carbowax" 4000
 - Sorbitol
 - "Dowicide" A

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The aqueous solution (about 10 kg) was spray-dried in a "Niro" utility spray drier using the above-described conditions and about 1 wt.% furned silica ("Cab sil" MS-7) was added to yield a free-fl wing powder having a residual wat r content of about 1%. About 2.3 kg of this powder was pelletized at ambient temperature in a California Pellet Mill, Model CL, using a die with 0.48 cm diameter orifices to yield small cylinders (pellets) having the following dimensions: diameter about 0.46 cm, length 0.33 to 0.9 cm (average about 0.6 cm), and a density of 1.28g/cc.

One hundred twenty g of the pellets (having a total surface area of about 1150 cm²) was placed in a phenolic resin impregnated cardboard sleeve (4.48 cm outside diameter, 4.25 cm inside diameter, 15.2 cm long). To retain the pellets in the sleeve as an aggregation, each end of the sleeve was fitted with a 4.3 cm diameter disc of "Scotch-Brite" fabric with the outer surface of the disc positioned about 0.5 cm from the sleeve end, and a bead of RTV silicone rubber was used to seal each disc at its juncture with the sleeve.

The loaded sleeve was inserted in the cartridge holder-nozzle assembly of Example 8 (Fig. 3) and connected to a hand portable, 2.5-gallon, water fire extinguisher. The extinguisher was filled with about 9.5 liters of 21°C tap water, pressurized with nitrogen gas to about 7 kgf/cm², and discharged over a period of 61 seconds.

An AFFF solution of good quality and fairly uniform concentration was produced as shown by the following data obtained on samples of foam collected at intervals during discharge.

		Table X	VII		
20	Time, Sec.	Refractive Index, n ₀ 20	Solute conc., g/l	Film speed, sec.	20
	2	1.3348	14.4	1.5	
	10	1.3346	12.8	1.5	
	20	1.3343	10.4	2	
25	30	1.3340	8.0	. 3	25
	.40	1.3338	6.4	6	
	50	1.3338	6.4	4	
	60	1.3338	6.4	5	
	Tap Water	1.3330	0	_	
30	Control	1.3340	8.0	. 4	30

After discharge, the cartridge was taken apart and the remaining pellets were dried in a circulating air oven at 110° for about 6 hours. The weight of the dried solids was 20.7 g, showing that 99.3 g had dissolved.

Example 10

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The following ingredients were separately pulverized and combined.

Table XVIII

Parts		
13.3	C ₆ F ₁₃ SO ₂ N(CH ₂ CHOHCH ₂ SO ₃)CH ₂ CH ₂ CH ₂ CH ₂ N ⁺ (CH ₃)CH ₂ CH ₂ OH	
6.7	C _B F ₁₇ SO ₃ K	
80	Sodium alkylsulfates ^{a.}	•

Solids obtained by evaporation of the water from "Polystep" B-25

Using a small laboratory extruder for plastics, the above powdered surfactant mixture was formed into continuous rod under heat (56—75°C.) and pressure. The pale amber rod (diameter 0.39 cm, density 1.41 g/cc) was cut into about 12.7 cm lengths, and 81 (total wt. 172.3 g) of these rods (having a total surface area of about 1275 cm²) were packed as an aggregation into a cardboard cartridge sleeve of the same type and dimensions as described in Example 9. The ends of the loaded sleeve were capped with "Scotch-Brite" fabric discs and sealed with RTV silicone adhesive as described in Example 9.

The loaded sleeve was inserted in the cartridge holder-nozzle assembly of Example 8 and connected to a 2.5-gallon, hand portable water fire extinguisher. The extinguisher was filled with about 9.5 liters of tap water, pressured to 7 kgf/cm² with nitrogen, and discharged completely over a period of 58.5 sec. An effective AFFF solution of quite uniform composition was produced as shown by the following data obtained on foam samples collected at intervals during discharge.

Tabl XIX 55 Time, Refractive Solute 55 Film speed, index, n₀20 sec. conc., g/l sec. 2 1.3334 . 3.2 24 10 1.3335 4.0 14 20 1.3335 4.0 12

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	Time,	Refractive	Solute	Film speed,	
	Sec.	Index n ₀ 20	conc., g/l	sec.	
	30	1.3335	4.0	12	
5	40	1.3335	4.0	13	5
•	50	1.3335	4.0	8	
	Tap Water	1.3330	0		
	Control	1.3340	8.0	7	
Th	e undissolved rods of the	cartridge weighed 12	7.3 a (after drying) indicating that 45.0 g of	f
	d dissolved.	outinago violgilos is		,	10

Example 11

After drying in a vacuum oven (75°C, 18 hrs) 557 g of the powdered surfactant product of Example 2 and 45.9 g of a powdered polysaccharide gum (K8A13) were combined and thoroughly mixed. This mixture was formed into a rod utilizing a small laboratory extruder at a barrel temperature of about 50°C. and a die temperature of about 65°C. The pale, amber rod (diameter 0.38 cm, density about 1.38 g/cc) was cut in about 1.3 cm lengths, and 120 g of these rod pieces (having a total surface area of about 1050 cm²) were placed as an aggregation into a sleeve of the same type and dimensions as described in Example 9. The ends of the loaded sleeve were capped with "Scotch-Brite" fabric discs and sealed as described in Example 9. The loaded sleeve was inserted in the cartridge holder-nozzle assembly of Example 8 and connected to a 2.5-gallon hand portable water fire extinguisher. The extinguisher was filled with about 9.5 liters of tap water (21°C), pressured to 7 kgf/cm² with nitrogen, and discharged completely over a period of 68 sec. An effective AFFF solution of quite uniform composition and properties was produced as shown by the following data obtained on foam samples collected at intervals during discharge.

collected	l at intervals during disch	arge.			
25	_	Table X	X		25
23	Time,	Refractive	Solute	Film speed,	
	sec.	index, n _D ²⁰	conc. g/l	sec.	
	2	1.3338	6.4	8	
	10	1.3338	6.4	7	
30	20	1.33375	6.0	10	30
30	30	1.3337	5.6	12	
	40	1.3337	5.6	12	
	50	1.3337	5.6	11	
	60	1.3337	5.6	13	
35	Tap Water	1.3330	0		35
J <i>U</i>	Control	1.3340	8.0	3	

The undissolved pellets of the cartridge weighed 57.7 g (after drying) indicating that 62.3 g had dissolved.

Various modifications and alterations will become apparent to those skilled in the art without departing from the scope and spirit of this invention.

Claims

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A hand portable fire extinguisher comprising a tank adapted to contain pressurized water, a valve surmounting said tank, a hose connected to said valve, a cartridge holder connected to said hose, an air aspirating nozzle connected to said cartridge holder, and a cartridge loaded in said cartridge holder and comprising a shaped body comprising a solid, coalesced mixture of water soluble fluoroaliphatic surfactant and water soluble fluorine-free surfactant, said shaped body having at least one exposed surface adapted to be contacted with water flowing through said cartridge holder to dissolve said shaped body and form an aqueous film-forming foam solution of relatively constant composition over the period of discharge of said water from said tank.

2. The extinguisher of claim 1, wherein said cartridge is a single shaped body comprising said mixture of surfactants.

3. The extinguisher of claim 1, wherein said cartridge comprises a plurality of said shaped bodies in the form of a water-permeable aggregation.

4. The extinguisher of claim 1, wherein said exposed surfac is adjacent to a water insoluble,
water permeable medium so as to provide said cartridge with at least one channel for said water to
pass therethrough in c intact with said surface.

5. The extinguisher of claim 1, wherein said cartridge comprises a layup comprising twe types of sheets in the form of a coil, one type of said sheet comprising said solid mixture and the other type of said sheet comprising a water insoluble, water permeable, open, resilient, three-dimensional web.

6. The extinguisher of claim 1, wherein said shaped body comprises a c mposite of said solid mixture distributed throughout a water insoluble, water permeable reinforcing matrix.

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7. The extinguisher of claim 1, wherein said fluoroaliphatic surfactant has the formula:

$(R_f)_n(Q)_m Z$

wherein R, is a fluorinated, saturated, monovalent, organic radical having a terminal perfluoromethyl group, containing from 3 to 20 carbon atoms, in which the carbon atoms of the chain are substituted 5 only by fluorine, chlorine or hydrogen atoms with no more than one hydrogen or chlorine atom for 5 every two carbon atoms, and in which a divalent oxygen or trivalent nitrogen atom, bonded only to carbon atoms, can be present in a skeletal chain, n is 1 or 2, Q is a multivalent linking group, m is an integer from 0 to 2, and Z is a water solubilizing polar group, and said fluorine-free surfactant is a synthetic, imputrescible, hydrocarbon-congrous, organic, fluorine-free surfactant water-soluble to at 10 least about 0.02 percent by weight in water at 25°C. and which substantially completely emulsifies at 10 least one phase of a mixture of equal volumes of cyclohexane and water at a concentration of about 0.1 to about 10 percent by weight of the water, the weight ratio of fluoroaliphatic surfactant to fluorine-free surfactant in said body being 10:1 to 1:25. 8. The extinguisher of claim 1, wherein said fluoroaliphatic surfactant comprises C₈F₁₇SO₃K and C₈F₁₃SO₂N(CH₂CHOHCH₂SO₃)CH₂CH₂CH₂CH₂CH₂CH₂CH₂OH, and said fluorine-free surfactant 15 comprises C₁₀H₂₁OSO₃Na and C₁₂H₂₅OSO₃Na. 9. The extinguisher of claim 1, wherein said solid mixture further comprises normally solid polyethylene glycol having a number average molecular weight in the range of about 1000 to 6000. 10. A hand portable fire extinguisher comprising a tank containing about 9.5 liters of water and about 2.8 liters of compressed gas at about 7 kgf/cm2, a valve and squeeze lever surmounting said 20 tank, a hose connected to said valve, a cartridge holder connected to said hose, an air aspirating nozzle connected to said cartridge holder, and a replaceable cartridge in said cartridge holder and comprising a layup comprising two types of sheets, one type of said sheet comprising a solid, coalesced mixture of water-soluble, fluoroaliphatic surfactant and water soluble, fluorine-free surfactant in the weight ratio 25 of 10:1 to 1:25 fluoroaliphatic surfactant to fluorine-free surfactant, the other type of said sheet 25 comprising a water insoluble, water permeable, open, resilient, three-dimensional web which permits contact of the surface of said one type of sheet with water flowing through said cartridge holder to dissolve said mixture and form an aqueous film-forming foam solution of relatively constant composition over a period of 45 to 90 seconds of discharge of said water from said tank. 11. A hand portable fire extinguisher comprising a tank containing about 9.5 liters of water and 30 about 2.8 liters of compressed gas at about 7 kgf/cm², a valve and squeeze lever surmounting said tank, a hose connected to said valve, a cartridge holder connected to said hose, an air aspirating nozzle connected to said cartridge holder, and a replaceable cartridge in said cartridge holder and comprising a water-permeable aggregation of a plurality of shaped bodies comprising a solid coalesced mixture of 35 water-soluble, fluoroaliphatic surfactant and water soluble, fluorine-free surfactant in the weight ratio 35 of 10:1 to 1:25 fluoroaliphatic surfactant to fluorine-free surfactant, said aggregation having exposed surface such that water flowing through said cartridge holder dissolves said mixture to form an aqueous film-forming foam solution of relatively constant composition over a period of 45 to 90 seconds of discharge of said water from sald tank. 12. A method of extinguishing a Class A or Class B fire, which comprises flowing a predetermined 40 amount of water under pressure in contact with the surface of at least one shaped body comprising a solid, coalesced, mixture of water soluble fluoroaliphatic surfactant and water soluble fluorine-free surfactant to produce an aqueous film-forming foam solution of relatively constant composition, and

applying said solution to said fire. 13. The method of claim 12, wherein said fire is a Class B fire of flammable liquid.

14. A shaped body comprising a solid, coalesced, mixture of water soluble fluoroaliphatic surfactant and water soluble, fluorine-free surfactant, said shaped body having at least one exposed surface which when contacted with a predetermined amount of flowing water is dissolved and forms an aqueous film-forming foam solution of relatively constant composition.

15. The shaped body of claim 14, comprising a composite comprising said solid mixture distributed throughout a water insoluble, water permeable reinforcing matrix.

16. The shaped body of claim 14 wherein said fluoroaliphatic surfactant has the formula

$(R_f)_n(Q)_m Z$

wherein R, is a fluorinated, saturated, monovalent, organic radical having a terminal perfluoromethyl 55 group, containing from 3 to 20 carbon atoms, in which the carbon atoms of the chain are substituted 55 only by fluorine, chlorine or hydrogen atoms with no more than one hydrogen or chlorine atom for every two carbon atoms, and in which a divalent xygen or trivalent nitrogen atom, b inded only to carbon atoms, can be pr sent in a skeletal chain, n is 1 or 2, Q is a multivalent linking group, m is an integer from 0 to 2, and Z is a water solubilizing polar gr up, and said fluorine-free surfactant is a 60 synthetic, imputrescible, hydrocarbon-congrous, organic, fluorine-free surfactant water-soluble to at 60 least about 0.02 percent by weight in wat rat 25°C, and which substantially completely emulsifies at

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least one phase of a mixture of equal volumes of cyclohexane and water at a concentration of about 0.1 to about 10 percent by weight of the water, the weight ratio of fluoroaliphatic surfactant to fluorine-free surfactant in said shaped body being 10:1 to 1:25.

17. The shaped body of claim 14 wherein said fluoroaliphatic surfactant comprises C₈F₁₇SO₃K and C₆F₁₃SO₂N(CH₂CHOHCH₂SO₃)CH₂CH₂CH₂CH₂CH₂CH₂CH₂OH, and said fluorine-free surfactant

comprises C₁₀H₂₁OSO₃Na and C₁₂H₂₅OSO₃Na.

18. The shaped body of claim 14, wherein said solid mixture further comprises normally solid polyethylene glycol having an average molecular weight in the range of about 1000 to 6000.

19. A water-permeable aggregation of a plurality of shaped bodies having predetermined exposed surface area each comprising a solid, coalesced mixture of water soluble, fluoroaliphatic surfactant and water soluble, fluorine-free surfactant, said aggregation which when contacted with a predetermined amount of flowing at a given rate water is dissolved and forms an aqueous film forming foam solution of relatively constant composition.

20. A cartridge comprising a sleeve containing the shaped body of claim 14 and a water insoluble, water permeable medium adjacent the surface of said shaped body so as to provide said cartridge with at least one channel for said water to pass therethrough in contact with said surface.

21. The cartridge of claim 20, further comprising a layup of two types of sheets, one type of said sheet comprising said solid mixture and the other type of sheet comprising a water insoluble, water permeable, open, resilient, three-dimensional web.

22. A cartridge comprising a water permeable aggregation of a plurality of shaped bodies of claim 20 19.

23. A cartridge comprising a sleeve containing a layup comprising two types of sheets, one type of sheet comprising a shaped body comprising a solid, coalesced, mixture of water soluble, fluoroaliphatic surfactant and water soluble, fluorine-free surfactant in the weight ratio of 10:1 to 1:25
25 fluoroaliphatic surfactant to fluorine-free surfactant, the other type of sheet comprising a water insoluble, water permeable, open, resilient, three-dimensional web which permits contact of the surface of sald one type of sheet with a predetermined amount of flowing water to dissolve said mixture and form an aqueous film-forming foam solution of relatively constant composition.

24. A cartridge holder loaded with the shaped body of claim 14.

25. A cartridge holder and air aspirating nozzle assembly, said cartridge holder being loaded with 30 at least one shaped body of claim 14.

26. A solid, coalesced, mixture of water soluble, fluoroaliphatic surfactant and water soluble, fluorine-free surfactant, said mixture upon being contacted with a stream predetermined amount of water flowing at a given rate dissolves to form an aqueous film-forming foam solution of relatively constant composition.

27. The mixture of claim 26 wherein said fluoroaliphatic surfactant has the formula:

$(R_f)_n(Q)_mZ$

wherein R_f is a fluorinated, saturated monovalent, organic radical having a terminal perfluoromethyl group, containing from 3 to 20 carbon atoms, in which the carbon atoms of the chain are substituted only by fluorine, chlorine or hydrogen atoms with no more than one hydrogen or chlorine atom for every two carbon atoms, and in which a divalent oxygen or trivalent nitrogen atom, bonded only to carbon atoms, can be present in a skeletal chain, n is 0 or 1, Q is a multivalent linking group, m is an integer from 0 to 2, and Z is a water solubilizing polar group, and said fluorine-free surfactant is a synthetic, imputrescible, hydrocarbon-congruous, organic, fluorine-free surfactant water-soluble to at least about 0.02 percent by weight in water at 25°C. and which substantially completely emulsifies at least one phase of a mixture of equal volumes of cyclohexane and water at a concentration of about 0.1 to about 10 percent by weight of the water, the weight ratio fluoroaliphatic surfactant to fluorine-free surfactant in said mixture being 10:1 to 1:25.

28. The mixture of claim 27, wherein said fluoroaliphatic surfactant comprises C_aF₁₇SO_aK and

28. The mixture of claim 27, wherein said fluoroaliphatic surfactant comprises C₈F₁₇SO₃K and C₈F₁₃SO₂N(CH₂CHOHCH₂SO₃)CH₂CH₂CH₂CH₂N⁺(CH₃)₂CH₂CH₂OH and said fluorine-free surfactant comprises C₁₀H₂₁OSO₃Na and C₁₂H₂₅OSO₃Na.

29. A hand portable fire extinguisher as claimed in Claim 1 substantially as herein described with

reference to any one of the Examples.

30. A hand portable fire extinguisher as claimed in Claim 1 substantially as herein described with reference to the acc impanying drawings.

31. A method f extinguishing fires as claimed in Claim 12 substantially as herein described.

32. A shaped body as claimed in Claim 13 substantially as herein described with reference to any one of the Examples.

33. A shaped body as claimed in Claim 13 substantially as herein described with reference to 60 Figure 3 of the accompanying drawings.

34. A cartridge as claimed in Claim 20 substantially as herein described with reference to any one of the Examples.

- 35. A cartridge as claimed in Claim 20 substantially as her in described with reference to Figure 4 of the accompanying drawings.
- 36. A solid coalesced mixture as claimed in Claim 26 substantially as herein d scribed with reference to any one of the Examples.

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ABSTRACT:

CHG DATE=19990617 STATUS=O> A hand portable fire extinguisher comprising a tank adapted to contain pressurized water, a valve surmounting

said tank, a

hose connected to said valve, a cartridge holder connected to said hose, an air

aspirating nozzle connected to said cartridge holder, and a cartridge loaded in

said cartridge holder and comprising a shaped body comprising a solid,

coalesced mixture of water soluble fluoroaliphatic surfactant and water soluble

fluorine-free surfactant, said shaped body having at least one exposed surface

adapted to be contacted with water flowing through said cartridge holder to

dissolve said shaped body and form an aqueous film-forming foam solution of

relatively constant composition over the period of discharge of said water from

said tank. Solid coalesced mixtures of water soluble fluorinated surfactant

and water soluble fluorine- free surfactant, shaped bodies comprising such

mixtures and cartridges containing a shaped body for use in the fire

extinguisher. <IMAGE>